PATENT COOPERATION TREATY

PCT

INTERNATIONAL SEARCH REPORT

(PCT Article 18 and Rules 43 and 44)

Applicant's or agent's file reference	FOR FURTHER ACTION	see Notification of Transmittal of International Search Report (Form PCT/ISA/220) as well as, where applicable, item 5 below.
P22412A/PKE/BOU		
International application No.	International filing date *(day/month/year)*	(Earliest) Priority Date *(day/month/year)*
PCT/GB 99/03331	07/10/1999	07/10/1998
Applicant		
GILTECH LIMITED et al.		

This International Search Report has been prepared by this International Searching Authority and is transmitted to the applicant according to Article 18. A copy is being transmitted to the International Bureau.

This International Search Report consists of a total of 3 sheets.

[X] It is also accompanied by a copy of each prior art document cited in this report.

1. **Basis of the report**

   a. With regard to the **language**, the international search was carried out on the basis of the international application in the language in which it was filed, unless otherwise indicated under this item.

   [ ] the international search was carried out on the basis of a translation of the international application furnished to this Authority (Rule 23.1(b)).

   b. With regard to any **nucleotide and/or amino acid sequence** disclosed in the international application, the international search was carried out on the basis of the sequence listing :

   [ ] contained in the international application in written form.

   [ ] filed together with the international application in computer readable form.

   [ ] furnished subsequently to this Authority in written form.

   [ ] furnished subsequently to this Authority in computer readble form.

   [ ] the statement that the subsequently furnished written sequence listing does not go beyond the disclosure in the international application as filed has been furnished.

   [ ] the statement that the information recorded in computer readable form is identical to the written sequence listing has been furnished

2. [ ] **Certain claims were found unsearchable** (See Box I).

3. [ ] **Unity of invention is lacking** (see Box II).

4. With regard to the **title,**

   [X] the text is approved as submitted by the applicant.

   [ ] the text has been established by this Authority to read as follows:

5. With regard to the **abstract,**

   [X] the text is approved as submitted by the applicant.

   [ ] the text has been established, according to Rule 38.2(b), by this Authority as it appears in Box III. The applicant may, within one month from the date of mailing of this international search report, submit comments to this Authority.

6. The figure of the **drawings** to be published with the abstract is Figure No. --

   [ ] as suggested by the applicant.

   [ ] because the applicant failed to suggest a figure.

   [ ] because this figure better characterizes the invention.

   [ ] None of the figures.

PATENT COOPERATION TREATY

REC'D 11 JAN 2001

WIPO PCT

PCT

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

(PCT Article 36 and Rule 70)

Applicant's or agent's file reference	FOR FURTHER ACTION	See Notification of Transmittal of International Preliminary Examination Report (Form PCT/IPEA/416)
P22412A/PKE/BOU		
International application No.	International filing date *(day/month/year)*	Priority date *(day/month/year)*
PCT/GB99/03331	07/10/1999	07/10/1998
International Patent Classification (IPC) or national classification and IPC A61K9/12		
Applicant GILTECH LIMITED et al.		

1. This international preliminary examination report has been prepared by this International Preliminary Examining Authority and is transmitted to the applicant according to Article 36.

2. This REPORT consists of a total of 6 sheets, including this cover sheet.

   ☐ This report is also accompanied by ANNEXES, i.e. sheets of the description, claims and/or drawings which have been amended and are the basis for this report and/or sheets containing rectifications made before this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions under the PCT).

   These annexes consist of a total of sheets.

3. This report contains indications relating to the following items:

   - I ☒ Basis of the report
   - II ☐ Priority
   - III ☐ Non-establishment of opinion with regard to novelty, inventive step and industrial applicability
   - IV ☐ Lack of unity of invention
   - V ☒ Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations suporting such statement
   - VI ☐ Certain documents cited
   - VII ☐ Certain defects in the international application
   - VIII ☒ Certain observations on the international application

Date of submission of the demand	Date of completion of this report
06/04/2000	08.01.2001
Name and mailing address of the international preliminary examining authority: European Patent Office D-80298 Munich Tel. +49 89 2399 - 0 Tx: 523656 epmu d Fax: +49 89 2399 - 4465	Authorized officer Hedegaard, A Telephone No. +49 89 2399 8644

Form PCT/IPEA/409 (cover sheet) (January 1994)

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No. PCT/GB99/03331

I. Basis of the r p rt

1. This report has been drawn on the basis of *(substitute sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to the report since they do not contain amendments (Rules 70.16 and 70.17).)*:
Description, pages:

1-27 as originally filed

Claims, No.:

1-24 as originally filed

2. With regard to the **language**, all the elements marked above were available or furnished to this Authority in the language in which the international application was filed, unless otherwise indicated under this item.

These elements were available or furnished to this Authority in the following language: , which is:

- ☐ the language of a translation furnished for the purposes of the international search (under Rule 23.1(b)).
- ☐ the language of publication of the international application (under Rule 48.3(b)).
- ☐ the language of a translation furnished for the purposes of international preliminary examination (under Rule 55.2 and/or 55.3).

3. With regard to any **nucleotide and/or amino acid sequence** disclosed in the international application, the international preliminary examination was carried out on the basis of the sequence listing:

- ☐ contained in the international application in written form.
- ☐ filed together with the international application in computer readable form.
- ☐ furnished subsequently to this Authority in written form.
- ☐ furnished subsequently to this Authority in computer readable form.
- ☐ The statement that the subsequently furnished written sequence listing does not go beyond the disclosure in the international application as filed has been furnished.
- ☐ The statement that the information recorded in computer readable form is identical to the written sequence listing has been furnished.

4. The amendments have resulted in the cancellation of:

- ☐ the description, pages:
- ☐ the claims, Nos.:
- ☐ the drawings, sheets:

5. ☐ This report has been established as if (some of) the amendments had not been made, since they have been considered to go beyond the disclosure as filed (Rule 70.2(c)):

(Any replacement sheet containing such amendments must be referred to under item 1 and annexed to this report.)

6. Additional observations, if necessary:

V. Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. Statement

Novelty (N)	Yes:	Claims	21-24
	No:	Claims	1-20
Inventive step (IS)	Yes:	Claims	
	No:	Claims	1-24
Industrial applicability (IA)	Yes:	Claims	1-24
	No:	Claims	

2. Citations and explanations
see separate sheet

VIII. Certain observations on the international application

The following observations on the clarity of the claims, description, and drawings or on the question whether the claims are fully supported by the description, are made:
see separate sheet

R Section V

Reasoned statement under Rule 66.2(a)(ii) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. Reference is made to the following documents:

   D1: WO-A-96 17595
   D2: EP-A-0 380 254
   D3: US-A-4 086 331
   D4: WO-A-94 00512
   D5: GB-A-1 503 897

   D1 discloses (see p. 11, l. 7-12 and example 1) formulations comprising a foamable gelling agent (e.g. alginate) and a slow-release precipitant therefor (calcium and silver ion releasing glass).

   D2 discloses (see claims 1-3 and example 1) formulations comprising a foamable gelling agent (e.g. alginate) and a precipitant therefor (di- or trivalent metal salt).

   D3 discloses (see claim 1 and example 1) formulations comprising a foamable gelling agent (gelatin) and a precipitant therefor (ferrous sulphate).

   D4 discloses (see claims 1, 5, 21, 24 and 25 and example 3) formulations comprising a foamable gelling agent (e.g. alginate) and a precipitant therefor (e.g. calcium carbonate).

   D5 discloses (see p. 2, l. 8-15 and p. 4, l. 108-110) formulations comprising a foamable gelling agent (carboxyethyl cellulose) and a precipitant therefor (trivalent metal ions).

2. The subject-matter of independent claims 1 and 12 is not novel (Art. 33(2) PCT) over D1-D5, each document taken separately (see above under item 1).

   It is here pointed out that neither the process step "wherein said slow-release

2. Contrary to the requirements of Rule 5.1(a)(ii) PCT, the relevant background art disclosed in the documents D2-D5 is not mentioned in the description, nor are these documents identified therein.

3. The description must be brought into conformity with the new claims to be filed; care should be taken during revision not to add subject-matter which extends beyond the content of the application as originally filed; Art. 34.2 (b) PCT.

   When amending the claims the Applicant is requested to identify those passages in the specification as originally filed on which the amended claims are based.

MUR-856445

PATENT COOPERATION TREATY

PCT

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

(PCT Article 36 and Rule 70)

Applicant's or agent's file reference	FOR FURTHER ACTION	See Notification of Transmittal of International Preliminary Examination Report (Form PCT/IPEA/416)
P22412A/PKE/BOU		
International application No.	International filing date *(day/month/year)*	Priority date *(day/month/year)*
PCT/GB99/03331	07/10/1999	07/10/1998
International Patent Classification (IPC) or national classification and IPC A61K9/12		
Applicant GILTECH LIMITED et al.		

1. This international preliminary examination report has been prepared by this International Preliminary Examining Authority and is transmitted to the applicant according to Article 36.

2. This REPORT consists of a total of 6 sheets, including this cover sheet.

   ☐ This report is also accompanied by ANNEXES, i.e. sheets of the description, claims and/or drawings which have been amended and are the basis for this report and/or sheets containing rectifications made before this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions under the PCT).

   These annexes consist of a total of sheets.

3. This report contains indications relating to the following items:

   - I ☒ Basis of the report
   - II ☐ Priority
   - III ☐ Non-establishment of opinion with regard to novelty, inventive step and industrial applicability
   - IV ☐ Lack of unity of invention
   - V ☒ Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations suporting such statement
   - VI ☐ Certain documents cited
   - VII ☐ Certain defects in the international application
   - VIII ☒ Certain observations on the international application

Date of submission of the demand	Date of completion of this report
06/04/2000	08.01.2001
Name and mailing address of the international preliminary examining authority: European Patent Office D-80298 Munich Tel. +49 89 2399 - 0 Tx: 523656 epmu d Fax: +49 89 2399 - 4465	Authorized officer Hedegaard, A Telephone No. +49 89 2399 8644

Form PCT/IPEA/409 (cover sheet) (January 1994)

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No. PCT/GB99/03331

I. Basis of the report

1. This report has been drawn on the basis of *(substitute sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to the report since they do not contain amendments (Rules 70.16 and 70.17).)*:
Description, pages:

1-27 as originally filed

Claims, No.:

1-24 as originally filed

2. With regard to the **language**, all the elements marked above were available or furnished to this Authority in the language in which the international application was filed, unless otherwise indicated under this item.

These elements were available or furnished to this Authority in the following language: , which is:

- ☐ the language of a translation furnished for the purposes of the international search (under Rule 23.1(b)).
- ☐ the language of publication of the international application (under Rule 48.3(b)).
- ☐ the language of a translation furnished for the purposes of international preliminary examination (under Rule 55.2 and/or 55.3).

3. With regard to any **nucleotide and/or amino acid sequence** disclosed in the international application, the international preliminary examination was carried out on the basis of the sequence listing:

- ☐ contained in the international application in written form.
- ☐ filed together with the international application in computer readable form.
- ☐ furnished subsequently to this Authority in written form.
- ☐ furnished subsequently to this Authority in computer readable form.
- ☐ The statement that the subsequently furnished written sequence listing does not go beyond the disclosure in the international application as filed has been furnished.
- ☐ The statement that the information recorded in computer readable form is identical to the written sequence listing has been furnished.

4. The amendments have resulted in the cancellation of:

- ☐ the description, pages:
- ☐ the claims, Nos.:
- ☐ the drawings, sheets:

5. ☐ This report has been established as if (some of) the amendments had not been made, since they have been considered to go beyond the disclosure as filed (Rule 70.2(c)):

(Any replacement sheet containing such amendments must be referred to under item 1 and annexed to this report.)

6. Additional observations, if necessary:

V. Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. Statement

Novelty (N)	Yes:	Claims	21-24
	No:	Claims	1-20
Inventive step (IS)	Yes:	Claims	
	No:	Claims	1-24
Industrial applicability (IA)	Yes:	Claims	1-24
	No:	Claims	

2. Citations and explanations
see separate sheet

VIII. Certain observations on the international application

The following observations on the clarity of the claims, description, and drawings or on the question whether the claims are fully supported by the description, are made:
see separate sheet

Re Section V

Reasoned statement under Rule 66.2(a)(ii) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. Reference is made to the following documents:

   D1: WO-A-96 17595
   D2: EP-A-0 380 254
   D3: US-A-4 086 331
   D4: WO-A-94 00512
   D5: GB-A-1 503 897

   D1 discloses (see p. 11, l. 7-12 and example 1) formulations comprising a foamable gelling agent (e.g. alginate) and a slow-release precipitant therefor (calcium and silver ion releasing glass).

   D2 discloses (see claims 1-3 and example 1) formulations comprising a foamable gelling agent (e.g. alginate) and a precipitant therefor (di- or trivalent metal salt).

   D3 discloses (see claim 1 and example 1) formulations comprising a foamable gelling agent (gelatin) and a precipitant therefor (ferrous sulphate).

   D4 discloses (see claims 1, 5, 21, 24 and 25 and example 3) formulations comprising a foamable gelling agent (e.g. alginate) and a precipitant therefor (e.g. calcium carbonate).

   D5 discloses (see p. 2, l. 8-15 and p. 4, l. 108-110) formulations comprising a foamable gelling agent (carboxyethyl cellulose) and a precipitant therefor (trivalent metal ions).

2. The subject-matter of independent claims 1 and 12 is not novel (Art. 33(2) PCT) over D1-D5, each document taken separately (see above under item 1).

   It is here pointed out that neither the process step "wherein said slow-release

precipitant is combined with said gelling agent during the foaming thereof" nor the intended use of the precipitant (as stabiliser) as defined in claim 1 can represent distinguishing features over D1-D5 since the claim as such is directed to a product.

3. The subject-matter of claims 21-24 is novel since a process as defined in claim 21 comprising the step of sterilising the dried foam by exposure to [SPEC0807]-irradiation or ethylene oxide has not been disclosed in the above-mentioned prior art documents.

4. With regard to the assessment of inventive step the documents D2 (see e.g. col. 8, l. 15-16), D3 (see e.g. col. 5, l. 3-4), D4 (see p. 12-13) and D5 (see p. 3, l. 91-106) have already disclosed the improved setting time and stability of foams made from formulations comprising foamable gelling agent and a precipitant therefor.

   Hence, it does not appear to represent any unexpected effect that the foams are stable enough to be sterilised as defined in present claim 21. Therefore, the subject-matter of the present application is not considered to involve an inventive step (Art. 33(3) PCT).

5. A positive international preliminary report for the subject-matter of the dependent claims 2-11, 13-20 and 22-24 can only be established when they refer to independent claims which meet the requirements of the PCT.

Re Section VIII

Certain observations on the international application

1. The term "or the like" used in claim 8 is vague and unclear and leaves the reader in doubt as to the meaning of the technical features to which it refers, thereby rendering the definition of the subject-matter of said claim unclear (Article 6 PCT).

2. Contrary to the requirements of Rule 5.1(a)(ii) PCT, the relevant background art disclosed in the documents D2-D5 is not mentioned in the description, nor are these documents identified therein.

3. The description must be brought into conformity with the new claims to be filed; care should be taken during revision not to add subject-matter which extends beyond the content of the application as originally filed; Art. 34.2 (b) PCT.

   When amending the claims the Applicant is requested to identify those passages in the specification as originally filed on which the amended claims are based.

PATENT COOPERATION TREATY

From the INTERNATIONAL BUREAU

PCT

NOTIFICATION OF ELECTION

(PCT Rule 61.2)

To:

Assistant Commissioner for Patents
United States Patent and Trademark Office
Box PCT
Washington, D.C.20231
ETATS-UNIS D'AMERIQUE

in its capacity as elected Office

Date of mailing (day/month/year) 01 May 2000 (01.05.00)	
International application No. PCT/GB99/03331	**Applicant's or agent's file reference** P22412A/PKE/BOU
International filing date (day/month/year) 07 October 1999 (07.10.99)	**Priority date** (day/month/year) 07 October 1998 (07.10.98)
Applicant GILCHRIST, Tom et al	

1. The designated Office is hereby notified of its election made:

[X] in the demand filed with the International Preliminary Examining Authority on:

06 April 2000 (06.04.00)

[ ] in a notice effecting later election filed with the International Bureau on:

2. The election [X] was

[ ] was not

made before the expiration of 19 months from the priority date or, where Rule 32 applies, within the time limit under Rule 32.2(b).

The International Bureau of WIPO **34, chemin des Colombettes** **1211 Geneva 20, Switzerland**	Authorized officer Anman QIU
Facsimile No.: (41-22) 740.14.35	Telephone No.: (41-22) 338.83.38

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under th PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece			TR	Turkey
BG	Bulgaria	HU	Hungary	ML	Mali	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MN	Mongolia	UA	Ukraine
BR	Brazil	IL	Israel	MR	Mauritania	UG	Uganda
BY	Belarus	IS	Iceland	MW	Malawi	US	United States of America
CA	Canada	IT	Italy	MX	Mexico	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NE	Niger	VN	Viet Nam
CG	Congo	KE	Kenya	NL	Netherlands	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NO	Norway	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	NZ	New Zealand		
CM	Cameroon			PL	Poland		
CN	China	KR	Republic of Korea	PT	Portugal		
CU	Cuba	KZ	Kazakstan	RO	Romania		
CZ	Czech Republic	LC	Saint Lucia	RU	Russian Federation		
DE	Germany	LI	Liechtenstein	SD	Sudan		
DK	Denmark	LK	Sri Lanka	SE	Sweden		
EE	Estonia	LR	Liberia	SG	Singapore		

FOAMABLE FORMULATION AND FOAM

The present invention is concerned with a foamable formulation and the foam formed therefrom.

A wide variety of gels, creams, ointments, lotions and other formulations are available for application to a body surface. The exact content of these compositions will vary depending upon the purpose of application. For example, a formulation may be applied to clean a body surface, to promote healing of any wound or injury, to prevent an exposed wound on the body from drying out, to prevent infection, etc. In certain circumstances the composition may include an active ingredient.

In our International Patent Application published 13 June 1996 under No WO-A-96/17595 we describe a foamable formulation which comprises a foamable carrier or gelling agent, for example an alginate gel, and an active ingredient, such as a water soluble glass powder.

The product described in WO-A-96/17595 represented a considerable advance over the use of gel or cream.

We have now found that by including a precipitant for the gelling agent, in a slow-release form within the composition, further improvements with regard to the setting time of the foam and its stability can be achieved. In particular, the added stability enables a pre-foamed pad to be sterilised by irradiation, ethylene oxide, or other conventional means.

Thus, the present invention provides a formulation comprising a foamed gelling agent combined with a slow-release precipitant therefor. The gelling agent may be any agent capable of forming a foam, although preferably the gelling agent is physiologically compatible and non-irritant when maintained in contact with the body surface. The gelling agent may be a gel, for example a sodium alginate gel, carageenan gel, sodium carboxymethylcellulose gel or mixtures thereof.

The precipitant is desirably intimately admixed throughout the whole of the foamed gelling agent, preferably during the foaming process. In certain circumstances however the presence of the precipitant on one surface of the foamed gelling agent may be sufficient to cause stabilisation of the foam. Examples of precipitants include stabilising crosslinking agents which render the gelling agent insoluble. Examples include salts of polyvalent metal ions such as calcium, zinc, copper, silver or aluminium as well as borates, glyoxal and amino-formaldehyde precondensates. In one embodiment, the polyvalent metal ion may be released from a water-soluble glass which is admixed into the foamable carrier in comminuted form. A copper ion-releasing water soluble glass, a zinc-ion releasing water soluble glass and mixtures thereof are particularly of interest.

The role of the precipitant is to stabilise the foamed gel so that a stable foam is produced. Generally, the stable foam should be produced within a reasonable time period since if the precipitant is too slow-acting, the foam structure will have collapsed prior to stabilisation. However, a very fast acting precipitant may not allow sufficient time for the gel to be foamed. Desirably, the precipitant stabilises the foamed gel over a time period of 1 minute to 120 minutes, preferably within 30 minutes, and most preferably within 15 minutes at ambient temperature. The foam is considered to be "cured" when it can be lifted and carefully handled without collapse. The solubility of the precipitant and hence the setting (cure) time of the foam may be varied by adjusting the pH of the composition, especially where the precipitant is based upon a calcium salt. Generally, the solubility of a calcium salt will be increased by lowering the pH. Typical pH adjusters include organic acids such as acetic, adipic, citric, fumaric, lactic, alginic and tartaric acids. Usually an amount of 0.5 g to 5 g of organic acid per 100 gel is sufficient. The organic acid may be admixed with the precipitant prior to foaming or, more preferably, may be admixed with the gelling agent prior to foaming.

Suitable precipitants include calcium citrate, calcium carbonate, calcium phosphate, calcium hydrogen phosphate ($CaHPO_4$), aluminium chloride, barium carbonate, barium phosphate, barium sulphate, barium chloride and zinc carbonate.

Where the gelling agent comprises an alginate gel, a carageenan gel or a carboxymethylcellulose gel one preferred precipitant is a calcium salt. Whilst calcium citrate has been used in the examples, other

slowly dissolving calcium salts are also suitable.

Where the gelling agent comprises carboxymethylcellulose gel one preferred precipitant is an aluminium salt.

In one embodiment the gelling agent and precipitant are packaged separately and only admixed during the foaming process or subsequent to foaming.

Alternatively, the precipitant may be included in a suspension (e.g. a suspension of calcium citrate and glycerine) which forms a separate layer on top of the gelling agent which remains substantially inert during handling and/or storage. Only once the operator desires to produce the foam, is the precipitant intimately admixed with the gelling agent (for example by shaking the container) and then promptly foamed. Using the precipitant in suspension form has the benefit that the suspension is easier to dispense from a pressurised container than a powder and also provides for more accurate dosing of unit precipitant per unit gelling agent.

Optionally, the formulation may comprise other additives such as decompactants which promote the desired foam structure or other foaming agents, plasticisers, humectants, preservatives, additives, sequestering agents or active ingredients such as antimicrobial agents, growth factors, hormones, living cells, etc.

The foam may be applied directly to the body area and allowed to produce a stable foam protective cover, for example over a wound. With the addition of the precipitants the cure of the foam is significantly

reduced, rendering the product more user friendly.

Alternatively, the foam can be produced onto a mould or other surface area, allowed to cure (for example by air drying or oven drying) and then applied to the body surface as a dressing. A foam sheet of this type is a preferred embodiment of the invention since it exhibits sufficient stability for easy handling whilst retaining a moist surface to promote wound healing. Optionally, the foam may be applied about a substrate (for example cloth, mesh, non-woven pad of alginate fibres, nylon, rayon, polylactid acid, polyglycolic acid, polycaprolactone or biocompatible glass fibres) which are then integrated into the foam pad produced.

As an example, the foam may be used to treat dermatological conditions (including psoriasis, atopic and allergic eczema). It may be convenient in this embodiment for the foam to deliver an active ingredient normally used to alleviate such conditions, for example a steroid such as hydrocortisone.

In another embodiment the foam may be used to treat burns or scalds, including sunburn.

In another embodiment the foam may be applied cosmetically, and for example may include skin moisturising agents, nutritional agents and growth factors suitable to promote skin regeneration. A foam intended for cosmetic use may include colorants or pigments so that the foam may be applied to the skin as a cosmetic or to disguise any blemishes in the skin.

The foam may be used prophylactically. In particular a foam containing a UV blocking agent may be applied to exposed areas of the skin to protect it from the

effects of the sun.

The formulation of the invention is applied to the body site of interest in the form of a foam and it is therefore essential that the composition undergoes a foaming process before application to the body. In the foaming process gas is forced into or is formed within the formulation to entrap small bubbles of gas therein, thereby forming the foam. Any suitably gas or gas producing system can be used to produce the foam. Mention may be made of butane and nitrous oxide, but other gases like air, nitrogen, hydrofluorocarbons such as HFC134a or 227, hydrocarbons like propane, isopropane or a mixture thereof, are also suitable. Conveniently the foam may be produced by conventional means such as by using aerosol technology.

The formulation according to the present invention may be stored in any convenient container until required. Generally, the container will be designed to preserve the sterile nature of the formulation. Conveniently the container will be provided with means to foam the composition when required. Details are given in WO-A-96/17595. A two can packaging and dispensing system, as described in our co-pending UK Patent Application No 9823029.5 (a copy of which is filed herewith), may be used to dispense the foam according to the present invention.

Generally, the foam will be produced from sterile ingredients.

Prior to the foaming process, the foamable carrier is preferably in the form of a gel. The gel may be sterilised and this is generally desirable where the foam is intended for medical use. Usually,

sterilisation will take place by autoclaving the formulation, since this is currently the most economic means of achieving sterilisation. Autoclaving at temperatures of from 100°C to 125°C for under ½ hour is normally sufficient. Generally, the autoclaving process should be as mild as possible, whilst being sufficient to sterilise the formulation. For example, autoclaving at temperatures of about 121°C for 15-20 minutes is acceptable. The autoclaved formulation may then be foamed when cool. It is also possible, however, to sterilise the formulation by other means, for example by γ-irradiation or e-beam irradiation. It has been found that autoclaving the gel may cause the MW of the foamable carrier to be slightly reduced. Consequently it may be desirable to select a foamable carrier having a higher MW than that ultimately required.

The foam forms an air-tight cover around any wound or injury to which it is applied, and this prevents that area from drying out and may also combat infection. The advantages of applying a topical product in the form of a foam include:

1. Easy rapid application,
2. Conforms to surface irregularities,
3. Insulates the wound,
4. Cools the tissues,
5. Offers antibacterial action to prevent infection,
6. Biocompatibility with tissue,
7. Suitable for use as a vehicle for the administration of pharmaceutical agents, and/or
8. Maintains a moist environment.

Generally, the formulation of the present invention will be applied directly to the body site of interest in the form of a foam, the foam being produced from any suitable device (such as an aerosol) immediately before application. It is, however, possible for a quantity of the foamed formulation to be produced and then applied onto the body site by any suitable means, for example by hand or by spatula. This method may be required for wounds having a narrow opening.

As stated above, the foam may also be produced on a suitable surface and then allowed to dry to produce a stable foam sheet which can be handled as described above without deterioration. Generally, the production of the sheet will take place under sterile conditions or may be sterilised after production. In the prior described foam product of WO-A-96/17595, it was not possible to provide a foamed pad product and then sterilise the pad by conventional means such as γ-irradiation, since it was found that the foam structure deteriorated during sterilisation. With the inclusion of the precipitant however, sterilisation of the pad is possible both by γ-irradiation, ethylene oxide sterilisation or other conventional means. This represents a very considerable advantage over the prior art product.

The foam sheet is generally produced by foaming the foamable carrier in the presence of the precipitant and allowing the foam to cure, usually by simply exposing the foam to the atmosphere to air dry at ambient temperature. Optionally the foam may be dried at elevated temperatures, for example may be oven dried. Desirably, the cure time of the foam is 40 minutes or less at ambient temperature and preferably the foam cures within 15 minutes, for example within 10 minutes.

Where the foam sheet is to be sterilised, it is advantageous to pre-treat the sheet prior to sterilisation in order to further stabilise the sheet. The difficulty with sterilising any foam of the type described is that the foam structure tends to deteriorate and collapse during the sterilisation process. The pre-treatment of the sheet preferably involves impregnating the sheet with further precipitant. Conveniently, this may entail immersing the sheet in a bath of the precipitant or of a solution of the precipitant. For example, the sheet may be immersed in a bath of calcium chloride or calcium citrate. To ensure that the precipitant penetrates into the centre of the foam sheet, the sheet may be gently squeezed whilst immersed in the bath. Generally, immersion of the sheet for a short period of time, such as 2 to 3 minutes, is sufficient. The sheet may then be removed from the bath of precipitant, washed in a mixture of de-ionised water and glycerine to enhance moisture content and then dried. The stabilised foam sheet may then be sterilised by gamma radiation or through use of ethylene oxide.

The ratio of de-ionised water : glycerine in the wash stage is preferably 19:1 by volume.

The treated foam sheet is desirably oven dried at relatively low temperatures, for example 100°C or less, preferably approximately 35°C.

In a preferred embodiment the foamable carrier includes a combination of copper and zinc ions, optionally in the form of water soluble glass(es). We have found that a foam containing appropriate quantities of these metal ions are particularly resistant to the deleterious effects of sterilisation. We hypothesise

that the copper and zinc ions act as scavenger of free radicals produced in the foam during sterilisation and which are, we believe, responsible for the breakdown in structure of the foam. Additionally, both copper and zinc ions have a radioprotective effect. Consequently, we consider that any material known for its use as a free radical scavenger and/or as a radioprotectant may likewise exhibit a protective effect on the foam structure during sterilisation.

Optionally the manufacture of a prefoamed product may envisage a continuous foaming process. The sheet may be divided into a convenient size and may be packaged. Optionally the foam sheet may be produced on contoured surface so that it is moulded to a pre-determined shape.

Examples of suitable foamable gelling agents for use in the composition of the present invention include (but are not limited to) alginate and derivatives thereof, carboxymethylcellulose and derivatives thereof, collagen, polysaccharides (including, for example, dextran, dextran derivatives, pectin, starch, modified starches such as starches having additional carboxyl and/or carboxamide groups and/or having hydrophillic side-chains, cellulose and derivatives thereof), agar and derivatives thereof (such as agar stabilised with polyacrylamide), carageenan, polyethylene oxides, glycol methacrylates, gelatin, gums such as xanthum, guar, karaya, gellan, arabic, tragacanth and locust bean gum. Also suitable are the salts of the aforementioned carriers, for example, sodium alginate. Mixtures of any of the aforementioned gelling agents may also be used, as required.

Preferred foamable gelling agents include alginate,

carageenan, carboxymethylcellulose, the derivatives and salts thereof and mixtures of any of these. Alginate (the derivatives or salts thereof, such as sodium and calcium alginate) are especially preferred. Foamable gelling agents having a molecular weight of from 10,000 to 200,000 kDa are preferred, especially over 100,000 kDa, for example 150,000 to 200,000 kDa, may be used.

The formulation may further comprise a foaming agent, which promotes the formation of the foam. Any agent having a surfactant character may be used. The surfactants may be cationic, non-ionic or anionic. Examples of suitable foaming agents include cetrimide, lecithin, soaps, silicones and the like. Commercially available surfactants such as Tween™ are also suitable. Cetrimide (which additionally has an anti-bacterial activity) is especially preferred.

The formulation of the present invention (and thus the foam) may be used to deliver pharmaceutically active agents, in particular to deliver such agents in a controlled release manner. Mention may be made of:

Antiseptics, Antibacterials and Antifungal agents, such as Chlorhexidine, acetic acid, polynoxylin, povidone iodine, mercurochrome phenoxyethanol, acridene, silver nitrate, dyes eg brilliant green, undecanoic acid, silver sulphadiazine, silver proteins and other silver compounds, metronidazole, benzaclonium chloride;

Nutritional agents, such as vitamins and proteins;

Growth factors and healing agents, including Ketanserin a serotonomic blocking agent;

Living Cells;

Enzymes include streptokinase and streptodormase;

Elements - zinc, selenium, cerium, copper, manganese, cobalt, boron, arsenic, chromium silver, gold, gallium;

Charcoal;

Desloughing and Debriding agents such as hypochlorite and hydrogen peroxide;

Astringents including potassium permanganate;

Antibiotics exemplified by neomycin and framycetin sulphate, sulfamylon, fusidic acid, mupirocin, bacitracin, gramicidin.

In addition the formulation of the present invention may further comprise other conventional additives such as plasticisers and humectants (such as glycerol, propane-1,2-diol, polypropylene glycol and other polyhydric alcohols), free radical scavengers to stabilise against the effects of sterilisation by irradiation, viscosity-adjusting agents, dyes and colorants, and the like.

Several experiments including comparatives tests have been made in order to demonstrate some of the advantages of the new compositions of the invention. Of course the embodiments described hereinbelow are submitted in order to better describe the invention and not to limit its scope.

EXAMPLE 1

PROCEDURE FOR MANUFACTURE OF UNIT BATCH (100 g) of ALGINATE GEL

Typically the alginate gels are made according to the following process:

1. De-ionised (DI) water is measured and poured into mixing vessel 1.
2. Desired amounts of suitable alginate (for example Keltone or Manucol) and glycerine are weighed using a calibrated balance, reading to 2 decimal places.
3. Alginate and glycerine are mixed together in a beaker until no lumps remain.
4. The whole alginate/glycerine mix is added very slowly to the water.
5. Once all the alginate/glycerine has been added to the water, the mixture is stirred until a smooth gel has formed.

Several different alginate gels have been made according the above process. They differ and are referred to by the amount of alginate (for example Keltone) used. For example the alginate gel code 6½ has the following composition:

GEL CODE	6½
DI Water	80 ml
Glycerine	25.22 g
Keltone	6.5 g
Unit Batch Wt	111.72 g

The above composition can be varied to include other

weights of alginate, which would be reflected in the gel code number. For example a composition having 8g alginate (plus 80ml DI water and 25.22g glycerine) would be designated gel code 8. Analogous gel codes are used when other gel formers (eg carageenan or CMC) are substituted for the alginate in the above composition.

In one embodiment, the gelling agent may be present in the form of a suspension, for example a suspension in glycerine. To avoid diluting the gelling agent, the gelling agent suspension may be made up with less glycerine such that the total quantity of glycerine present in the gelling agent mixture and in the precipitant suspension adds up to the required amount. For example, the glycerine in the gelling agent mixture and precipitant suspension may be varied as follows:

Glycerine per 80 ml DI water and 6 g alginate (g)	**Glycerine in precipitant suspension (g)**
25.22	0
23.0	2.22
20.0	5.22
18.22	7.0
15.0	10.22

The above is illustrated with respect to a gel code 6 composition, but the division of glycerine may be made for other gel code compositions, and is also not limited to the specific volumes illustrated above.

PROCEDURE FOR FOAM PRODUCTION

The propellant used to produce the foam can be compressed gases such as air, nitrogen, nitrous oxide or air, hydrofluorocarbons such HFC134a or 227 or hydrocarbons including propane, isopropane, n-butane, isobutane and 2-methylbutane.

Propellant vapour pressure can range from 0 to 110 PSIG at 70°C although the preferred range is 20 to 70 PSIG. Values within this range can be achieved for example by blending the three hydrocarbons propane, isobutane and butane. Calor Aerosol Propellants (CAP) sold by Calor Gas Ltd Slough may be used as propellant gas, when a blend of propane, isobutane and butane is used the proportions can be as follows:

Grade	Propane %	Isobutane %	n Butane%
CAP 30	11	29	60
CAP 40	22	24	54
CAP 70	55	15	30

A foam according to the invention can advantageously be produced following the following process:

1. 100 g of a gel according to the invention is poured to an aerosol canister.
2. 2.5 g of calcium citrate (food grade) is added to the canister.
3. A valve is crimped onto the canister.
4. Air is purged from the canister.
5. 4.5 g of propellant gas is added into the canister (65:35 CAP 40 : Isopentane propellant) and an actuator is positioned on the valve.
6. The canister is shaken vigorously for 20-30 seconds.

7. The canister is inverted and the foam dispensed.

EXAMPLE 2

Using a range of water-based gel formulations detailed below tests were done to improve the "setting" time and stability of the gel and its foam.

Preferred alginate compositions have an amount of alginate ranging from 5-9g in the composition set out in Example 1. Preferred alginates are Keltone HV and Manucol DMF.

Experiment 1. Gel Code 6½ Alginate gel and foam mixed with calcium citrate compared to Gel Code 6½ alginate gel alone

Foamed gel with calcium citrate

2.5 g calcium citrate was added to 100 g of gel and the foamed gel was spread out onto plastic sheeting. The resultant foam pad was liftable in 15 minutes.

Foamed gel without calcium citrate

The above experiment was reproduced by foaming the gel on its own as described above. The "setting" time of the foam was 10 hours.

The experiments were repeated using 100 g unfoamed gel with and without calcium citrate. Similar setting times to those observed for the foamed gels were obtained (15 minutes and 10 hours respectively) before the gel pads were liftable.

Conclusion: Calcium citrate speeds up and controls the setting time of the gel and the foam.

Experiment 2. Gel Code 8 Alginate gel mixed with water

soluble glass (WSG) containing phosphate and boron compared to gel code 8 alginate gel alone.

The WSG was comprised as follows:
28.5M% CaO
3M% Ag
5M% B_2O_3
18.5M% MgO
45M% P_2O_5

Foamed gel with WSG

2.5 g of WSG was mixed with 100 g gel and the foamed mixture was spread out onto plastic sheeting. The resultant foam pad was liftable in 120 mins.

Foamed gel without WSG

The above experiment was repeated by foaming the gel on its own. The "setting" time of the foam was approximately 10 hours.

The experiments were repeated using 100 g unfoamed gel with and without WSG. Similar setting times to those observed for the foamed gels were obtained (120 minutes and 10 hours respectively) before the gel pads were liftable.

Conclusion: WSG speeds up and controls the setting time of the gel and the foam.

Experiment 3. Gel Code 4 Carageenan gel mixed with calcium citrate compared to gel code 4 gel alone

Foamed gel with calcium citrate

3 g of calcium citrate was mixed with 100 g gel and the foamed mix was spread out onto plastic sheeting. The resultant foam pad was liftable in 120 mins.

Foamed gel without calcium citrate

The above experiment was repeated by foaming gel on its own as described above. The "setting" time of the foam was 10 hours.

The experiments were repeated using 100 g unfoamed gel with and without calcium citrate. Similar setting times to those observed for the foamed gels were obtained (120 minutes and 10 hours respectively) before the gel pads were liftable.

Experiment 4. Gel Code 4½ Carageenan gel and gel code 6½ alginate gel mixed with calcium citrate compared to gel code 4½ carageenan gel and gel code 6½ alginate gel alone

Foamed gel with calcium citrate

2.5 g of calcium citrate was mixed with (50 g alginate and 50 g carageenan) gel and the foamed mix was spread out onto plastic sheeting. The resultant foam pad was liftable in 15 mins.

Foamed gel without calcium citrate

The above experiment was repeated by foaming the mixed gel on its own. The "setting" time of the foam pad was 10 hours.

The experiments were repeated using 100 g unfoamed gel with and without calcium citrate. Similar setting times to these observed for the foamed gels were obtained (120 minutes and 10 hours respectively) before the gel pads were liftable.

Experiment 5. Gel Code 6½ Alginate gel mixed with calcium citrate and added bentone IPM gel

2.5 g calcium citrate was added to 100 g of gel with 1g bentone IPM gel, admixed in an aerosol canister and dispensed therefrom as a foam onto a plastic surface. The resultant foam pad was liftable in 12 minutes. Bentone IPM gel is an admixture of isopropyl myristate, sterealkonium hectorite and propylene carbonate.

Conclusion: Calcium citrate and bentone gel control the setting time of the foam. Bentone gel also acts as a reological agent and assists in the smoothness of delivery from the can.

Experiment 6. Gel Code 6½ Alginate gel mixed with calcium citrate and added cetrimide

2.5 g calcium citrate was added to 100 g of alginate gel with 1g cetrimide in an aerosol canister and foamed onto a plastic surface. The resultant foam pad was liftable in 15 minutes.

Conclusion: Calcium citrate speeds up the setting time of the foam. Cetrimide increases the cell structure of the product.

Experiment 7. Gel Code 6½ Alginate gel mixed with calcium citrate and added Tween 20

2.5 g Calcium citrate was added to 100 g of alginate gel with 1g Tween 20 and foamed onto a plastic surface. The resultant foam pad was liftable in 12 minutes.

Conclusion: Calcium citrate speeds up the setting time of the gel. The additive Tween 20 gave a much smoother delivery and an airier foam. Tween 80, 60 and 40 were also tried and all assisted in the delivery and product cell structure.

Experiment 8. Gel Code 4 Carboxmethyl cellulose and gel code 6½ alginate gel mixed with calcium citrate compared to the gel alone

2.5 g calcium citrate was added to (50 g CMC & 50 g alginate gel) and then the mixture was foamed onto a plastic surface. The resultant foam pad was liftable in 25 minutes. The gel foamed on its own was liftable overnight (approx. 10 hours).

Experiment 9. Gel Code 4 Carboxmethyl cellulose gel mixed with aluminium chloride compared with the gel alone

2 g aluminium chloride was mixed with 100 g CMC gel. The gel was spread onto a plastic surface. The resultant gel was liftable instantly. The gel alone was liftable overnight (approx. 10 hours).

Experiment 10. Gel Code 6 Alginate gel mixed with citric acid compared to gel code 6 alginate gel alone

2.5 g of citric acid was mixed with 100 g alginate gel and the mix was spread out onto plastic sheeting. The resultant gel pad was liftable in 120 mins. 100 g of the gel alone was spread onto plastic sheeting and the resultant pad was only liftable overnight (approx. 10 hours).

Experiment 11. Gel Code 6½ Alginate gel was mixed with the following powders on a 100 g gel: 2.5 g powder basis

Powder	Results as a gel	Results as a foam
Calcium Chloride	Gel pad was formed instantly	Fast setting foam
Calcium Sulphate	Gel pad formed reasonably quickly	Foam set reasonably quickly
Aluminium Chloride	Gel pad formed instantly	Fast setting foam
Calcium Metaborate	Gel pad formed instantly	Fast setting foam

Experiment 12. Setting performances of a foam of a gel code 6½ alginate gel as a function of the amounts of calcium citrate.

Batch No	Amount of calcium citrate per 100 g gel	Result
DM02 210798	4 g	Not dispensed - set in can
DM03 210798	3 g	Very difficult to dispense. 9½ minutes to set.
DM04 210798	2.5 g	Easier to dispense than above. 18½ minutes to set
DM05 210798	2.25 g	Taking longer to set. 20 minutes.
DM02 200798	2 g	Setting time - 40 minutes

Experiment 13. Gel Code 6½ alginate gel with calcium citrate and isopentane.

100g gel code 6½ alginate gel was admixed with varying amounts of calcium citrate (2 to 4g), added to isopentane and mixed thoroughly before being spread onto a glass sheet. The isopentane vaporises at ambient temperatures and boils off through the gel leaving a foam pad of similar consistency to those produced by dispersion from an aerosol can. After half-an-hour the foam pads were liftable.

EXAMPLE 3

A. Gel code 5 alginate gel mixed with calcium citrate

The gel was prepared by mixing together alginate (5g Keltone HV), 20g glycerine and 80ml de-ionised water. 5.22g glycerine was then added to 2.5g calcium citrate and a suspension of precipitant was created. The resultant gel and the suspension of precipitant were added to an aerosol can and a valve fitted. The can was purged of air, filled with 4.5g CAP 40 butane, shaken and dispensed. The foam produced was well mixed and set in 15 minutes.

B. Gel code 5 alginate gel mixed with calcium citrate

Experiment A was repeated using the same weight of Manucol LKX (5g) instead of Keltone HV. The resultant foam set within 12 minutes.

C. Gel code 5 alginate gel mixed with calcium citrate

The gel was prepared by mixing together alginate (5g Keltone HV), 20g glycerine and 80ml de-ionised water.

5.22g glycerine was then added to 2.5g calcium citrate and a suspension of precipitant was created. The resultant gel was added to the bottom can of the two can packaging system (see our co-pending UK Patent Application No 9823029.5) and the suspension or precipitant was added to the top can. The cans were prepared in the usual way. The two can packaging system was activated and the foam was dispensed. The foam produced was well mixed and set in 15 minutes.

D. Gel code 5 alginate gel mixed with calcium citrate

Experiment C was repeated using the same weight of Manucol LKX instead of Keltone HV. The resultant foam set within 12 minutes.

The set foam from A, B, C and D were then further processed by first immersing the foam in a solution of 2.5% calcium chloride solution for 2 minutes, rinsing in de-ionised water and then finally rinsing in a 1% glycerine solution. The foam pads were then dried in the oven at 35°C and packaged in sterilisable pouches.

The resultant sterilised pads were compared with can reference 2 below (see Example 4). The foams produced in the two can system had a more even pore size throughout compared to those made in a one can system. Comparing the suspension with the powder/gel mix showed no difference in the structure of the final product.

EXAMPLE 4

A 1 litre batch of gel code 5 alginate gel was manufactured. Nine bottom cans of a two can packaging system as described in our co-pending UK Patent Application No 9823029.5 were filled with 100g gel in

each. Nine top cans were made up with varying powders as detailed below. The cans were prepared in their usual way. The two can packaging system was activated and the foam was dispensed.

Once cured the foams were processed by varying a) the concentration of the calcium chloride immersion solution and b) the final wash concentration of the glycerine solution. All samples were halved and then oven dried at 40°C. The first half sample was removed after 8 hours and the second half after 16 hours. Once the foam pads had been processed they were packaged in EtO sterilisable airtight packaging as soon as they came out of the oven. The samples were sent for EtO sterilisation and examined on their return.

Can Ref	Top Can Component	Ca Chloride Conc.	Glycerine Sol Conc.	Drying Time	Description of Alginate Pad After EtO Sterilisation
1	2.5 g Ca Citrate	1%	1%	8 hrs	Flexible, soft & sponge-like
				16 hrs	Flexible, soft & sponge-like
2	2.5 g Ca Citrate	2.5%	1%	8 hrs	Moist, flexible & sponge-like
				16 hrs	Flexible, soft & sponge-like
3	2.5 g Ca Citrate	5%	1%	8 hrs	Dry pad with limited flexibility
				16 hrs	Dry pad with limited flexibility
4	2.5 g Ca Citrate	2.5%	2%	8 hrs	Moist, flexible, soft & sponge-like
				16 hrs	Moist, flexible, soft & sponge-like
5	2.5 g Ca Citrate	2.5%	2.5%	8 hrs	Moist, flexible, sponge-like pad
				16 hrs	Moist, flexible, sponge-like pad
6	2.5 g Ca Citrate	2.5%	5%	8 hrs	Moist, flexible, soft & sponge-like
				16 hrs	Moist, flexible, soft & sponge-like
7	2 g Ca Citrate 2 g Activated Charcoal	2.5%	5%	8 hrs	Moist, flexible, soft & sponge-like
				16 hrs	Moist, flexible, soft & sponge-like
8	2 g Ca Citrate 2 g Cu/Zn WSG	2.5%	5%	8 hrs	Moist, flexible, soft & sponge-like
				16 hrs	Moist, flexible, soft & sponge-like
9	2.5 g Ca Citrate 2 g Povidone Iodine	2.5%	5%	8 hrs	Moist, flexible, soft & sponge-like
				16 hrs	Moist, flexible, soft & sponge-like

EXAMPLE 5

Experiment A

A 600 g batch of gel code 5 was made up using Manucol DMF as the gelling agent. This batch was split into six equal parts and inserted into the bottom can of a dual can aerosol system. The top cans were made up containing 1.5 g calcium citrate and varying amounts of alginic acid (½ g increments from 0 to 2½ g). Once preparation was complete the cans were foamed out simultaneously and the setting time for each foam was recorded.

Can Number	Gel Weight	Calcium Citrate Weight	Alginic Acid Weight	Setting Time
1	100 g	1.5 g	0 g	20 mins
2	100 g	1.5 g	0.5 g	16 mins
3	100 g	1.5 g	1.0 g	14 mins
4	100 g	1.5 g	1.5 g	10 mins
5	100 g	1.5 g	2.0 g	9 mins
6	100 g	1.5 g	2.5 g	8 mins

Experiment B

Three 100 g batches of gel code 5 was made up using Manucol DMF as the gelling agent with alginic acid incorporated (0 g, 1 g and 2 g added). Each batch was filled into bottom cans and top cans were made up containing 1.5 g calcium citrate. Once preparation complete the cans were foamed out simultaneously and the setting times for each can recorded.

Can Number	Gel Weight	Calcium Citrate Weight	Alginic Acid Weight	Setting Time
7	100 g	1.5 g	1 g	8 mins
8	100 g	1.5 g	2 g	6 mins
9	100 g	1.5 g	0 g	20 mins

CLAIMS

1. A physiologically acceptable formulation for application to a body as a foam, said formulation comprising a foamable gelling agent and a slow-release precipitant therefor, wherein said slow-release precipitant is combined with said gelling agent during the foaming thereof and stabilises the foamed form of the gelling agent.

2. A formulation as claimed in Claim 1 wherein said precipitant is packaged separately to said gelling agent prior to foaming.

3. A formulation as claimed in either one of Claims 1 and 2 wherein said gelling agent is alginate, carboxymethylcellulose, collagen, a polysaccharide, agar, a polyethylene oxide, a glycol methacrylate, gelatin, a gum, or salts or derivatives of any of these, or mixtures thereof.

4. A formulation as claimed in Claim 3 wherein said gelling agent is alginate, carboxymethyl-cellulose, carageenan gel, the derivatives or salts thereof, or mixtures thereof.

5. A formulation as claimed in any one of Claims 1 to 4, wherein said gelling agent has a molecular weight of from 10,000 to 200,000 kDa.

6. A formulation as claimed in any one of Claims 1 to 5, wherein said precipitant is a salt of calcium, zinc, copper, silver or aluminium; borates; glyoxal; or amino-formaldehyde pre-condensates

7. A formulation as claimed in any one of Claims 1 to 6 further containing a foaming agent.

8. A formulation as claimed in Claim 7 wherein said foaming agent is cetrimide, lecithin, a soap, silicone, a surfactant or the like.

9. A formulation as claimed in any one of Claims 1 to 8 wherein the gelling agent comprises an alginate gel, a carageenan gel or a carboxymethylcellulose gel and wherein the precipitant is a calcium salt.

10. A formulation as claimed in any one of Claims 1 to 8 wherein the gelling agent comprises carboxymethylcellulose gel and wherein the precipitant is an aluminium salt.

11. A formulation as claimed in any one of Claims 1 to 10 further comprising an organic acid in an amount of 0.5 g to 5.0 g per 100 g gelling agent.

12. A physiologically acceptable foam comprising a foamed gelling agent stabilised by a precipitant.

13. The foam as claimed in Claim 12 in the form of a cured foam sheet.

14. A foam as claimed in Claim 12 wherein said precipitant is packaged separately to said gelling agent prior to foaming.

15. A foam as claimed in any one of Claims 12 to 14 wherein said gelling agent is alginate, carboxymethylcellulose, collagen, a polysaccharide, agar, a polyethylene oxide, a glycol methacrylate, gelatin, a gum, or salts or

derivatives of any of these, or mixtures thereof.

16. A foam as claimed in Claim 15 wherein said gelling agent is alginate, carboxymethyl- cellulose, carageenan gel, the derivatives or salts thereof, or mixtures thereof.

17. A foam as claimed in any one of Claims 12 to 16, wherein said gelling agent has a molecular weight of from 10,000 to 200,000 kDa.

18. A foam as claimed in any one of Claims 12 to 17, wherein said precipitant is a salt of calcium, zinc, copper, silver or aluminium; borates; glyoxal; or amino-formaldehyde pre-condensates

19. A foam as claimed in any one of Claims 12 to 18 further containing a foaming agent.

20. A foam as claimed in Claim 19 wherein said foaming agent is cetrimide, lecithin, a soap, silicone, a surfactant or the like.

21. A process of sterilising a foam for medical or veterinary use, said process comprising:

a) foaming a formulation of Claims 1 to 11 and allowing said foamed formulation to cure;

b) treating said foam with precipitant;

c) optionally, washing said treated foam;

d) drying said treated form; and

e) sterilising said dried foam by exposure to γ-irradiation or ethylene oxide.

22. The process of Claim 21 wherein said treated foam is washed in a de-ionised water/glycerine mixture prior to drying.

23. The process of either one of Claims 21 and 22 wherein the treated foam is oven dried at temperatures below 100°C.

24. The process of any one of Claims 21 to 23 wherein the foam is immersed in a bath of calcium chloride or calcium citrate solution as precipitant.

INTERNATIONAL SEARCH REPORT

International Application No
PCT/GB 99/03331

A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 A61K9/12

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 7 A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 96 17595 A (GILTECH LTD ;GILCHRIST THOMAS (GB); GILCHRIST EILIDH (GB)) 13 June 1996 (1996-06-13) cited in the application page 3, line 17 -page 4, line 15; claims 1-10; example 1	1-9, 11-20
A	page 8, line 5 -page 10, line 17 page 11, line 7 - line 12 ---	21-24
X	EP 0 380 254 A (MINNESOTA MINING & MFG) 1 August 1990 (1990-08-01) column 4, line 19 -column 5, line 25; claims; examples --- -/--	1-6,9, 11,12, 14-18

[X] Further documents are listed in the continuation of box C.

[X] Patent family members are listed in annex.

° Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search	Date of mailing of the international search report
20 January 2000	27/01/2000

Name and mailing address of the ISA	Authorized officer
European Patent Office, P.B. 5818 Patentlaan 2 NL – 2280 HV Rijswijk Tel. (+31–70) 340–2040, Tx. 31 651 epo nl, Fax: (+31–70) 340–3016	Marttin, E

Form PCT/ISA/210 (second sheet) (July 1992)

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category °	Citation of document, with indication,where appropriate, of the relevant passages	Relevant to claim No.
X	US 4 086 331 A (NEUMANN PETER) 25 April 1978 (1978-04-25)	1,3,7,8, 12,14, 15,19,20
A	column 1, line 35 - line 48; claims; examples 1,2 column 2, line 4 - line 23 column 2, line 63 -column 3, line 8	6,18
X	WO 94 00512 A (ALBANY INT CORP ;BAKIS GEORGE (US); EAGLES DANA BURTON (US); HAGOO) 6 January 1994 (1994-01-06) page 6, line 9 - line 13; claims 1,5,11-25; example 3 page 7, line 17 - line 24 page 8, line 10 -page 9, line 10 page 16, line 1 - line 2	1,3-9, 12,13, 15-20
A	example 4	21,24
X	GB 1 503 897 A (BAYER AG) 15 March 1978 (1978-03-15) page 2, line 8 - line 15; example 1 page 2, line 127 -page 3, line 20 page 4, line 31 - line 44 page 4, line 108 - line 117	1,3,4,6, 10

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No
PCT/GB 99/03331

Patent document cited in search report		Publication date	Patent family member(s)		Publication date
WO 9617595	A	13-06-1996	AU	3990095 A	26-06-1996
			CA	2203367 A	13-06-1996
			EP	0797430 A	01-10-1997
			JP	10509724 T	22-09-1998
EP 0380254	A	01-08-1990	US	5057606 A	15-10-1991
			CA	2006882 A	24-07-1990
			DE	69010221 D	04-08-1994
			DE	69010221 T	26-01-1995
			JP	2232234 A	14-09-1990
			US	5089606 A	18-02-1992
US 4086331	A	25-04-1978	NONE		
WO 9400512	A	06-01-1994	AU	672214 B	26-09-1996
			AU	4543793 A	24-01-1994
			BR	9305552 A	20-12-1994
			CA	2116037 A	06-01-1994
			EP	0612331 A	31-08-1994
			FI	940735 A	16-02-1994
			JP	2644626 B	25-08-1997
			JP	6510330 T	17-11-1994
			NO	940546 A	17-02-1994
			US	5840777 A	24-11-1998
			US	5851461 A	22-12-1998
GB 1503897	A	15-03-1978	DE	2544777 A	21-04-1977
			BE	847025 A	07-04-1977
			BR	7606678 A	16-11-1977
			DK	450976 A	08-04-1977
			ES	452143 A	01-10-1977
			FR	2326937 A	06-05-1977
			IE	43740 B	20-05-1981
			IT	1076815 B	27-04-1985
			JP	52047926 A	16-04-1977
			LU	75938 A	25-05-1977
			NL	7611041 A	13-04-1977

INTERNATIONAL SEARCH REPORT

ational Application No
PCT/GB 99/03331

A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 A61K9/12

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 7 A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X ✓	WO 96 17595 A (GILTECH LTD ;GILCHRIST THOMAS (GB); GILCHRIST EILIDH (GB)) 13 June 1996 (1996-06-13) cited in the application page 3, line 17 -page 4, line 15; claims 1-10; example 1	1-9, 11-20
A	page 8, line 5 -page 10, line 17 page 11, line 7 - line 12	21-24
X ✓	EP 0 380 254 A (MINNESOTA MINING & MFG) 1 August 1990 (1990-08-01) column 4, line 19 -column 5, line 25; claims; examples	1-6,9, 11,12, 14-18

-/--

[X] Further documents are listed in the continuation of box C.

[X] Patent family members are listed in annex.

° Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search	Date of mailing of the international search report
20 January 2000	27/01/2000

Name and mailing address of the ISA	Authorized officer
European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016	Marttin, E

Form PCT/ISA/210 (second sheet) (July 1992)

INTERNATIONAL SEARCH REPORT

International Application No
PCT/GB 99/03331

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X ✓	US 4 086 331 A (NEUMANN PETER) 25 April 1978 (1978-04-25)	1,3,7,8, 12,14, 15,19,20
A	column 1, line 35 - line 48; claims; examples 1,2 column 2, line 4 - line 23 column 2, line 63 -column 3, line 8	6,18
X ✓	WO 94 00512 A (ALBANY INT CORP ;BAKIS GEORGE (US); EAGLES DANA BURTON (US); HAGOO) 6 January 1994 (1994-01-06) page 6, line 9 - line 13; claims 1,5,11-25; example 3 page 7, line 17 - line 24 page 8, line 10 -page 9, line 10 page 16, line 1 - line 2	1,3-9, 12,13, 15-20
A	example 4	21,24
X ✓	GB 1 503 897 A (BAYER AG) 15 March 1978 (1978-03-15) page 2, line 8 - line 15; example 1 page 2, line 127 -page 3, line 20 page 4, line 31 - line 44 page 4, line 108 - line 117	1,3,4,6, 10

Form PCT/ISA/210 (continuation of second sheet) (July 1992)

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No
PCT/GB 99/03331

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9617595 A	13-06-1996	AU 3990095 A	26-06-1996
		CA 2203367 A	13-06-1996
		EP 0797430 A	01-10-1997
		JP 10509724 T	22-09-1998
EP 0380254 A	01-08-1990	US 5057606 A	15-10-1991
		CA 2006882 A	24-07-1990
		DE 69010221 D	04-08-1994
		DE 69010221 T	26-01-1995
		JP 2232234 A	14-09-1990
		US 5089606 A	18-02-1992
US 4086331 A	25-04-1978	NONE	
WO 9400512 A	06-01-1994	AU 672214 B	26-09-1996
		AU 4543793 A	24-01-1994
		BR 9305552 A	20-12-1994
		CA 2116037 A	06-01-1994
		EP 0612331 A	31-08-1994
		FI 940735 A	16-02-1994
		JP 2644626 B	25-08-1997
		JP 6510330 T	17-11-1994
		NO 940546 A	17-02-1994
		US 5840777 A	24-11-1998
		US 5851461 A	22-12-1998
GB 1503897 A	15-03-1978	DE 2544777 A	21-04-1977
		BE 847025 A	07-04-1977
		BR 7606678 A	16-11-1977
		DK 450976 A	08-04-1977
		ES 452143 A	01-10-1977
		FR 2326937 A	06-05-1977
		IE 43740 B	20-05-1981
		IT 1076815 B	27-04-1985
		JP 52047926 A	16-04-1977
		LU 75938 A	25-05-1977
		NL 7611041 A	13-04-1977

US 0976398304P1

Creation date: 01-16-2004
Indexing Officer: KKHAMBAY - KHOUTHONG KHAMBAY
Team: OIPEBackFileIndexing
Dossier: 09763983

Legal Date: 04-11-2001

No.	Doccode	Number of pages
1	M903	1

Total number of pages: 1

Remarks:

Order of re-scan issued on .....